## AMENDMENTS TO THE CLAIMS

 (currently amended) A method for making a cell-matrix construct for use as a heart valve comprising

implanting into an animal a cell-matrix construct comprising a fibrous matrix in the shape of a heart valve or heart valve leaflet, wherein the matrix is formed of a biocompatible, biodegradable polymer having seeded therein cells selected from the group consisting of endothelial cells, myofibroblasts, skeletal muscle cells, vascular smooth muscle cells, myocytes, fibromyoblasts, and ectodermal cells,

wherein the sell-matrix construct can withstand repeated stress and strain synthetic

biodegradable polymer provides the biomechanical properties of a heart valve or leaflet until the

seeded cells can lay down their own extracellular matrix, and

the matrix is formed so that the cells attach to and proliferate on it to the edges of the matrix.

- (currently amended) The method of claim 1 wherein the matrix is seeded with dissociated parenehymal-or connective tissue cells.
- (previously presented) The method of claim 1 wherein the matrix is first cultured at a
  first site in a patient prior to being transplanted to a second site.
- (previously presented) The method of claim 1 wherein the matrix is in the form of a heart valve leaflet.

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5. (previously presented) The method of claim 1 wherein the cell-matrix construct is seeded with vascular smooth muscle cells and endothelial cells and implanted to form a heart

valve.

6-8. (cancelled)

9. (previously presented) The method of claim 1 wherein the cell-matrix construct is

formed of a polymer selected from the group consisting of poly(lactide) (PLA), poly(glycolic

acid) (PGA), poly(lactide-co-glycolide) (PLGA), poly(caprolactone), polycarbonates,

polyamides, polyamiydrides, polyamino acids, polyortho esters, polyacetals, polycyanoacrylates,

and degradable polyurethanes.

10. (cancelled)

11. (previously presented) The method of claim 1 wherein the cell-matrix construct

contains interconnected pores in the range of between approximately 100 and 300 microns.

12. (previously presented) The method of claim 1 wherein the cell-matrix construct

includes growth factors.

13. (previously presented) The method of claim 12 wherein the growth factors are

selected from the group consisting of heparin binding growth factor (hbgf), transforming growth

factor alpha or beta (TGF), alpha fibroblastic growth factor (FGF), epidermal growth factor

(TGF), vascular endothelium growth factor (VEGF), insulin, glucagon, estrogen, nerve growth

factor (NGF) and muscle morphogenic factor (MMP).

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14. (previously presented) The method of claim 1 wherein the cell-matrix further comprises bioactive factors incorporated to between one and 30% by weight.

15-17. (cancelled)

18. (new) The method of claim 1 wherein the cell-matrix is first cultured in a bioreactor to form a fibrous tissue-polymeric construct before implantation.

19. (new) The method of claim 18 wherein the bioreactor is an animal.

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